

United States Patent and Trademark Office

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS / P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NO. FILING DATE FIRST NAMED INVENTOR ATTORNEY DOCKET NO. CONFIRMATION NO. 02/11/2000 06501-056001 09/502,698 Shin-Ichi Funahashi 5541 **EXAMINER** 26161 01/14/2005 FISH & RICHARDSON PC MERTZ, PREMA MARIA 225 FRANKLIN ST PAPER NUMBER ART UNIT BOSTON, MA 02110

DATE MAILED: 01/14/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)
Office Action Summary	09/502,698	FUNAHASHI ET AL.
	Examiner	Art Unit
	Prema M Mertz	1646
The MAILING DATE of this communicated Period for Reply	ation appears on the cover sheet wit	h the correspondence address
A SHORTENED STATUTORY PERIOD FOI THE MAILING DATE OF THIS COMMUNIC. - Extensions of time may be available under the provisions of after SIX (6) MONTHS from the mailing date of this commun - If the period for reply specified above is less than thirty (30) of - If NO period for reply is specified above, the maximum statul - Failure to reply within the set or extended period for reply will - Any reply received by the Office later than three months after - earned patent term adjustment. See 37 CFR 1.704(b).	ATION. 37 CFR 1.136(a). In no event, however, may a relication. days, a reply within the statutory minimum of thirty tory period will apply and will expire SIX (6) MONIII, by statute, cause the application to become ABA	ply be timely filed (30) days will be considered timely. HS from the mailing date of this communication. ANDONED (35 U.S.C. § 133).
Status		
1)⊠ Responsive to communication(s) filed	on <u>21 October 2004</u> .	
2a) This action is FINAL . 2b)⊠ This action is non-final.	
3)☐ Since this application is in condition fo closed in accordance with the practice	,	•
Disposition of Claims		
4) ⊠ Claim(s) 3 and 9-37 is/are pending in to 4a) Of the above claim(s) 9-34 is/are with 5) □ Claim(s) is/are allowed. 6) ⊠ Claim(s) 3, 35-37 is/are rejected. 7) □ Claim(s) is/are objected to. 8) □ Claim(s) are subject to restriction	rithdrawn from consideration.	
Application Papers		
9) The specification is objected to by the E 10) The drawing(s) filed on is/are: a Applicant may not request that any objection Replacement drawing sheet(s) including the	a) accepted or b) objected to bon to the drawing(s) be held in abeyand	ce. See 37 CFR 1.85(a).
11) The oath or declaration is objected to b		· · · · · ·
Priority under 35 U.S.C. § 119		
12) Acknowledgment is made of a claim for a) All b) Some * c) None of: 1. Certified copies of the priority do 2. Certified copies of the priority do 3. Copies of the certified copies of application from the Internationa * See the attached detailed Office action for	ocuments have been received. Ocuments have been received in Ap the priority documents have been of the Bureau (PCT Rule 17.2(a)).	plication No received in this National Stage
Attachment(s)		
1) Notice of References Cited (PTO-892)	4) Interview Su	
 Notice of Draftsperson's Patent Drawing Review (PTC3) Information Disclosure Statement(s) (PTO-1449 or PT Paper No(s)/Mail Date 		/Mail Date ormal Patent Application (PTO-152) -

Art Unit: 1646

DETAILED ACTION

- 1. A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 10/21/2004 has been entered.
- 2. Claims 1-2, 4-5 have been canceled. Claims 9-34 have been withdrawn from consideration as drawn to a non-elected invention. Previously presented claims 3, 35-37 are pending in the instant application and are under consideration by the Examiner.
- 3. Receipt of applicant's arguments and amendments filed on 10/21/2004 is acknowledged.
- 4. Applicant's arguments filed on 4/29/04 have been fully considered and were non-persuasive. The remaining issues are stated below.
- 5. The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

Claim Rejections - 35 USC § § 101/112, first paragraph

6. Claims 3, 35-37 are rejected under 35 U.S.C. 101.

This rejection is maintained for reasons of record set forth at pages 3-5 of the previous Office action (4/18/03) and pages 2-3 of the Office action (9/22/03).

Applicants argue that the polypeptides at issue can be used for generating antibodies to be employed in detecting liver cells and lung cancer tissues, and that this use is specific, substantial and credible and have cited Example 12 of the USPTO Utility

Art Unit: 1646

Guidelines Training material in this regard. Furthermore, Applicants argue that post-filing date evidence discloses that the claimed polypeptides bind to 5-HT_{2C} receptors which are known to be involved in neural transmission and have cited the Ullmer et al publication demonstrating a 454 amino acid polypeptide fragment that is 98% identical to a portion of SEQ ID NO:1 and 2 that binds to the serotonin 5-HT_{2C} receptor. However, contrary to Applicants arguments, it is clear from the instant specification that the instantly claimed protein is what is termed an "orphan protein" in the art. As shown in Figure 2 of the instant specification, there is only 98% homology between a "portion" of the instant protein (from amino acid 921-1373) and the receptor protein of Ullmer et al. However, the Ullmer et al publication discloses that the MUPP1 protein which is a member of the PDZ protein family interacts with the C-terminus of the 5-HT_{2C} receptors i.e. the PDZ domain protein MUPP1 is a scaffolding protein that interacts with the 5-HT_{2C} receptor. This disclosure in the post-filing reference was not disclosed in the present application as filed.

Applicant has also traversed this rejection on the premise that a claimed protein can be employed in identifying compounds that agonize or antagonize activity of 5-HT_{2C} receptor, which utility Applicants argue is a credible, specific and substantial utility. Applicants argue that the methods for screening for compounds and for generating antibodies is a specific, substantial and credible utility. However, the employment of a protein of the instant invention, in identifying compounds that agonize or antagonize activity of the 5-HT_{2C} receptor is not a credible, substantial or specific utility. To grant Applicants a patent encompassing an isolated protein of as yet undetermined biological significance would be to grant Applicant a monopoly "the metes and bounds" of which

Art Unit: 1646

"are not capable of precise delineation". That monopoly "may engross a vast, unknown, and perhaps unknowable area" and "confer power to block off whole areas of scientific development, without compensating benefit to the public" *Brenner v. Manson*, *Ibid*). To grant Applicant a patent on the claimed polypeptide based solely upon an assertion that a portion of the instant protein has 98% homology to the 454 amino acid fragment of Ullmer et al, is clearly prohibited by this judicial precedent since the compensation to the public is not commensurate with the monopoly granted. The instant protein has no demonstrated function.

Applicants also argue that the claimed polypeptide has a specific and substantial utility because it can be used in identification of certain tissues including liver and lung tumor cells and this supports the utility of the antibodies for detection of either liver cells or lung cancer cells. However, the employment of the claimed polypeptide in such a method is not a substantial or specific utility, because the instant polypeptide has not been shown to be differentially expressed in normal and lung tumors. Applicants have failed to show differential expression of the instant nucleic acid in normal lung tissue and in lung tumor tissue. Applicant is not being required to identify a ligand for protein, and a physiological process mediated thereby and a disease or disorder for which that protein is a marker. Applicant is only required to identify one substantial credible utility and, as stated in the previous office action, the employment of this protein only as the subject of further research does not satisfy the utility requirement of 35 U.S.C. § 101 because the courts have interpreted this statute as requiring an invention to have "substantial utility" "where specific benefit exists in currently available form". The employment of a protein of the instant invention, as a tissue specific marker is not a substantial or specific utility.

Art Unit: 1646

All human proteins can invariably be classified into two categories, those which are expressed in a tissue or developmentally specific manner and those which are expressed ubiquitously. It can be alleged that any protein which is expressed in a tissue specific manner can be employed to detect the tissue in which it is expressed in a sample. Alternately, a human protein which is expressed ubiquitously can be employed to detect the presence of any human tissue in a sample. Such utilities are analogous to the assertion that a particular protein can be employed as a molecular weight marker, which is neither a specific or substantial utility.

The examiner must simply provide sound reasoning in support of a conclusion that an element is lacking from a specification, and this has been done. In the instant case, it is the responsibility of Applicant to disclose a specific utility for the claimed invention and factually unsupported assertions like those presented i.e. in detection of lung tumors, are not specific utilities on their face that they need not be "proven" wrong. The following is an excerpt from M.P.E.P. 2138.05:

Utility for the invention must be known at the time of the reduction to practice. Wiesner v. Weigert, 212 USPQ 721, 726 (CCPA 1981) (except for plant and design inventions); Azar v. Burns, 188 USPQ 601, 604 (Bd. Pat. Inter. 1975) (a composition and a method cannot be actually reduced to practice unless the composition and the product produced by the method have a practical utility); Ciric v. Flanigen, 185 USPQ 103, 105 - 6 (CCPA 1975) ("when a count does not recite any particular utility, evidence establishing a substantial utility for any purpose is sufficient to prove a reduction to practice"; "the demonstrated similarity of ion exchange and adsorptive properties between the newly discovered zeolites and known crystalline zeolites ... have established utility for the zeolites of the count"); Engelhardt v. Judd, 151 USPQ 732, 735 (CCPA 1966) (When considering an actual reduction to practice as a bar to patentability for claims to compounds, it is sufficient to successfully demonstrate utility of the compounds in animals for somewhat different pharmaceutical purposes than those asserted in the specification for

Art Unit: 1646

humans.); Rey - Bellet v. Engelhardt,181 USPQ 453, 455 (CCPA 1974) (Two categories of tests on laboratory animals have been considered adequate to show utility and reduction to practice: first, tests carried out to prove utility in humans where there is a satisfactory correlation between humans and animals, and second, tests carried out to prove utility for treating animals.).

A PROBABLE UTILITY MAY NOT BE SUFFICIENT TO ESTABLISH UTILITY

A probable utility does not establish a practical utility, which is established by actual testing or where the utility can be "foretold with certainty." Bindra v. Kelly, 206 USPQ 570, 575 (Bd. Pat. Inter. 1979) (Reduction to practice was not established for an intermediate useful in the preparation of a second intermediate with a known utility in the preparation of a pharmaceutical. The record established there was a high degree of probability of a successful preparation because one skilled in the art may have been motivated, in the sense of 35 U.S.C. 103, to prepare the second intermediate from the first inter mediate. However, a strong probability of utility is not sufficient to establish practical utility.); Wu v. Jucker, 167 USPQ 467, 472 (Bd. Pat. Inter. 1968) (screening test where there was an indication of possible utility is insufficient to establish practical utility). But see Nelson v. Bowler, 206 USPQ 881, 885 (CCPA 1980) (Relevant evidence is judged as a whole for its persuasiveness in linking observed properties to suggested uses. Reasonable correlation between the two is sufficient for an actual reduction to practice)."

Furthermore, use of the protein to raise antibodies is analogous to the assertion that a particular DNA can be employed as a molecular weight marker, which is neither a specific or substantial utility.

There has to be physiological significance for the polypeptide disclosed in the specification. This requirement is analogous to basic scientific characterization, however, in the instant case no substantial benefit for the claimed protein is currently disclosed, but an exploratory significance. In conclusion, Applicants arguments with respect to utility of the instant polypeptide, are found to be non-persuasive. Contrary to Applicants arguments, the instant specification does not disclose a single credible,

Art Unit: 1646

specific or substantial utility for the instant polypeptide. The initial burden to demonstrate or present such is on Applicants.

Claims 3, 35-37 also remain rejected under 35 U.S.C. 112, first paragraph, as failing to adequately teach how to use the instant invention. Claims 3, 35-37 stand rejected under 35 U.S.C. § 112, first paragraph, because the instant specification does not teach how to use the invention for those reasons of record in pages 3-5 of the previous Office action (4/18/03) and pages 2-3 of the Office action (9/22/03).

Claim rejections-35 USC § 112, second paragraph

7. Claims 3, 35-37 are rejected under 35 U.S.C. § 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 3 is unclear because it recites "comprising the sequence of SEQ ID NO:1" rather than "comprising the amino acid sequence of SEQ ID NO:1".

Claim 35 is unclear because it recites "comprising SEQ ID NO:2" rather than "comprising the amino acid sequence of SEQ ID NO:2".

Claim 36 is unclear because it recites "consists of SEQ ID NO:2" rather than "consists of the amino acid sequence of SEQ ID NO:1".

Claim 37 is unclear because it recites "consists of SEQ ID NO:2" rather than "consists of the amino acid sequence of SEQ ID NO:2".

Conclusion

No claim is allowed.

Advisory Information

Art Unit: 1646

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Prema Mertz whose telephone number is (571) 272-0876. The examiner can normally be reached on Monday-Friday from 7:00AM to 3:30PM (Eastern time).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Anthony Caputa, can be reached on (571) 272-0829.

Official papers filed by fax should be directed to (571) 273-8300. Faxed draft or informal communications with the examiner should be directed to (571) 273-0876.

Information regarding the status of an application may be obtained from the Patent application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Prema Mertz Ph.D.
Primary Examiner
Art Unit 1646
January 11, 2005